

Standard Reference Material® 1632c

Trace Elements in Coal

(Bituminous)

This Standard Reference Material (SRM) is intended primarily for use in the evaluation of techniques employed in the analysis of coals and materials of a similar matrix. SRM 1632c consists of 50 g of bituminous coal ground to pass a 250 µm (60 mesh) sieve, homogenized, and bottled under an argon atmosphere.

Certified Values: The certified concentrations for 15 elements, expressed as mass fractions [1] on a dry basis, are provided in Table 1. The certified values for all elements, except sulfur, are based on two or more critically evaluated independent methods as described by Schiller and Eberhardt [2]. The certified value for sulfur is based on a single NIST primary method, isotope dilution thermal ionization mass spectrometry (ID-TIMS) [3]. A NIST certified value is a value for which NIST has the highest confidence in its accuracy in that all known or suspected sources of bias have been investigated or accounted for by NIST [4].

Reference Values: The reference values for 26 constituents, expressed as mass fractions on a dry basis, are provided in Table 2. The reference values for carbon, nitrogen, and fluorine are based on two independent methods. The reference value for ash content is based on results from 64 laboratories participating in an interlaboratory study in conjunction with the Canada Centre for Mineral and Energy Technology (CANMET) Service Program for the Evaluation of Codes and Standards (CANSPECS) using ASTM methods [5,6]. The reference values for other constituents are from a single NIST analytical method. Reference values are noncertified values that are the best estimate of the true value; however, the values do not meet NIST criteria for certification and are provided with associated uncertainties that may reflect only measurement precision and may not include all sources of uncertainty [4].

Information Values: Information values for 13 selected elements, volatile matter content [6,7], and gross calorific value are provided in Table 3 for information purposes only. These are noncertified values with no uncertainty assessed. Summary statistics reported by CANSPECS for SRM 1632c, which was included as an unknown in the CANSPECS 54 Coal and CANSPECS 54 Ash Round Robins, are provided in the addendum to this certificate to demonstrate user experience with this material using conventional methods and to better characterize the matrix. The ash used in the CANSPECS 54 Ash Round Robin was prepared from the coal used to produce SRM 1632c and ashed in accordance with ASTM D 6357-00 [8]. The CANSPECS interlaboratory results should **NOT** be used as substitutes for NIST values.

Expiration of Certification: The certification of SRM 1632c is valid, within the measurement uncertainties specified, until **01 May 2010**, provided the SRM is handled in accordance with the instructions given in this certificate (see "Instructions for Use"). This certification is nullified if the SRM is contaminated or otherwise modified.

Maintenance of SRM Certification: NIST will monitor representative samples of this SRM over the period of its certification. If substantive changes occur that affect the certification before the expiration of this certificate, NIST will notify the purchaser. Registration (see attached sheet) will facilitate notification.

The coordination of the technical measurements leading to certification was performed by R.R. Greenberg of the NIST Analytical Chemistry Division.

Stephen A. Wise, Chief Analytical Chemistry Division

Robert L. Watters, Jr., Chief Measurement Services Division

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Statistical analyses leading to certified and reference values were performed by J.H. Yen and D.D. Leber of the NIST Statistical Engineering Division.

The support aspects involved in the preparation of this SRM were coordinated through the NIST Measurement Services Division.

The coal for this SRM was donated by Consol Coal Sales¹, Inc. (Pittsburgh, PA).

INSTRUCTIONS FOR USE

Sampling: The SRM should be thoroughly mixed by rotating the bottle before sampling. A minimum sample mass of 250 mg should be used for analytical determinations to be related to elemental concentration values provided. The calorific value and ash content were determined using a minimum sample mass of 1 g. The SRM should be stored in its original, tightly sealed bottle away from sunlight and intense sources of radiation.

Drying: In order to relate measurements to the certified and reference values that are expressed on a dry mass basis, users should determine a drying correction at the time of each analysis. The correction is determined by oven drying a separate 1 g sample in a nitrogen atmosphere at $107~^{\circ}\text{C} \pm 3~^{\circ}\text{C}$ to a constant mass [6]. During drying at NIST, the mass loss of SRM 1632c samples was observed to stabilize between 77 minutes and 91 minutes [9]. The average mass loss measured at NIST for SRM 1632c was 2.02 % (1 s = 0.13 %, n = 12). Oven drying at $105~^{\circ}\text{C}$ for 2 hours in an air atmosphere and desiccator drying for 5 days over fresh magnesium perchlorate are also acceptable drying methods. No significant difference between these two methods was observed. Moisture content determined by vacuum drying for 2 hours at ambient temperatures was found to be lower than oven or desiccator drying by about $0.5~^{\circ}$ to $1~^{\circ}$ absolute.

A NIST study was also conducted to quantify the difference between drying in air and nitrogen atmospheres for SRM 1632c. For the same time and temperature conditions, the average mass loss for oven drying the SRM in an air atmosphere was 1.97 % (1 s = 0.11 %, n = 12).

SOURCE, PREPARATION, AND ANALYSIS¹

Source and Preparation of Material: The coal for this SRM was obtained from the Bailey Mine of the Consol Coal Company in southwestern Green County, PA. This mine produces bituminous coal obtained from the Pittsburgh seam which is considered to be one of the most extensively mined and economically important coal seams in the United States. The collection of the approximately 340 kg of washed coal was performed under the direction of L.W. Rosendale, Consol Coal Research and Development. The coal was air dried and subsequently pulverized to pass a 250 µm (60 mesh) sieve by a company under contract to NIST. At NIST, the entire lot was divided using the spinning riffle technique into two portions. One portion of the lot was sealed for long term storage in foil bags filled under an argon atmosphere. The remaining portion was further divided by the spinning riffler technique and bottled under an argon atmosphere. Several hundred bottles of the SRM 1632c lot were labeled SRM 1630a *Trace Mercury in Coal* and issued on 11 February 1999.

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¹Certain commercial equipment, instruments, or materials are identified in this certificate in order to adequately specify the experimental procedure. Such identification does not imply recommendation or endorsement by the National Institute of Standards and Technology, nor does it imply that the materials or equipment identified are necessarily the best available for the purpose.

Table 1. Certified Mass Fractions for Selected Elements (Dry basis) in SRM 1632c

Major Constituents

Minor Constituents

Elements	Mass Fra	action	Elements	Mass (Frac %)	etion
Hydrogen Sulfur	5.11 ± 1.462 ±	0.12 0.051	Potassium Chlorine	0.1100 0.1139		0.0033 0.0041
		Trace Elements				
Elements	Mass Fra (mg/k		Elements	Mass (m	Frac g/kg	
Antimony	0.461 ±	0.029	Selenium	1.326	±	0.071
Barium	41.1 ±	1.6	Sodium	298.8	±	4.8
Cobalt	$3.48 \pm$	0.20	Strontium	63.8	\pm	1.4
Manganese	$13.04 \pm$	0.53	Thorium	1.40	\pm	0.03
Mercury	$0.0938 \pm$	0.0037	Zinc	12.1	±	1.3
Rubidium	7.52 \pm	0.33				

The uncertainty in the certified value for sulfur is expressed as an expanded uncertainty, U, and is calculated according to the method described in the ISO and NIST Guides [10]. The observed sulfur variation was much greater than expected for the analytical technique used. Therefore, a prediction interval was used to account for the sulfur variability in this material. The expanded uncertainty is calculated as $U = ku_c$, where u_c is intended to represent, at the level of one standard deviation, the combined effect of uncertainty components associated with the measurement uncertainty and sulfur inhomogeneity, and k is a coverage factor. The coverage factor, k = 2.3, is determined from the Student's t-distribution with 8 degrees of freedom and corresponds to a 95 % prediction interval [11].

The uncertainties for all other certified values are weighted means of results from two or more analytical methods. For these certified values and reference values, the uncertainty is calculated as $U = ku_c + B$. The quantity u_c is the combined standard uncertainty calculated according to the ISO and NIST Guides [10], which accounts for the combined effect of the within variance for all methods at one standard deviation. The coverage factor, k, is determined from the Student's t-distribution corresponding to the appropriate associated degrees of freedom and 95 % confidence for each element. The term, B, is a bias adjustment for the difference between methods, which is the maximum difference between the certified value and the method means [2].

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Table 2. Reference Values (Dry basis) for SRM 1632c

Major Constituents

Minor Constituents

Elements	Mass F		Elements	Mass Fraction (%)
Carbon (Total) Nitrogen Silicon		0.25 0.06 0.034	Aluminum Calcium Iron Magnesium	$\begin{array}{cccc} 0.915 & \pm & 0.0137 \\ 0.145 & \pm & 0.030 \\ 0.735 & \pm & 0.011 \\ 0.0384 & \pm & 0.0032 \end{array}$
		Trace Elements		
Elements	Mass F	- ** * * - *	Elements	Mass Fraction (mg/kg)
Arsenic Boron Bromine Cadmium Cerium Chromium Cesium Copper Europium	6.18 ± 62 ± 18.7 ± 0.072 ± 11.9 ± 13.73 ± 0.594 ± 6.01 ± 0.1238 ±	2 0.4 0.007 0.2 0.20 0.010 0.25 0.0033	Fluorine Hafnium Lead Nickel Samarium Scandium Titanium Uranium Vanadium	$\begin{array}{cccccccccccccccccccccccccccccccccccc$
	Mass F			
Ash	7.16 ±	0.05		

The uncertainty in the carbon, nitrogen, and ash content reference values is calculated as an expanded uncertainty, $U = ku_c + B$. The quantity u_c is intended to represent, at the level of one standard deviation, the combined effect of within laboratory measurement uncertainty and between laboratory uncertainty [10]. The coverage factor, k, is determined from the Student's t-distribution with the appropriate associated degrees of freedom. The term, k, is a bias adjustment for the difference between methods, which is the maximum difference between the reference value and the method means [2].

The uncertainty in the reference value for fluorine is expressed as an expanded uncertainty, $U = ku_c$, calculated according to the methods in the ISO and NIST Guides [10]. The quantity u_c represents, at the level of one standard deviation, the combined uncertainty due to the potential effects within and between laboratories in the assessment of fluorine. The quantity k is the coverage factor used to specify the approximate confidence level of the expanded uncertainty interval. The value of the coverage factor, k = 3.182, is determined from the Student's t-distribution with 3 effective degrees of freedom and a confidence level of 95 %.

The uncertainties for all other reference values are the expanded uncertainties calculated according to the ISO and NIST Guides [10].

Supplemental Information: The information values given in Table 3 are not certified and are given as additional information on the matrix. **NOTE:** Gross calorific value and volatile matter content [6,7] may decrease with time due to sample degradation.

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Table 3. Information Values (Dry basis) for SRM 1632c

Elements	Mass Fraction (mg/kg)	Elements	Mass Fraction (mg/kg)
Beryllium	1.0	Silver	0.1
Bismuth	0.1	Tellurium	0.05
Gallium	3	Thallium	0.4
Germanium	5	Tin	1
Lithium	8	Yttrium	4
Molybdenum	0.8	Zirconium	16
Niobium	1		

Volatile Matter Content 36.0 %

Gross Calorific Value $32.10 \text{ MJ} \cdot \text{kg}^{-1} (13\ 802\ \text{Btu}_{\text{th}} \cdot \text{lb}^{-1})$

Table 4. Methods of Analysis

Element	Method	Element	Method
Aluminum	INAA	Magnesium	INAA
Antimony	ICP-MS, INAA	Manganese	INAA, ICP-MS
Arsenic	INAA	Mercury	ID-CV-ICP-MS, RNAA
Ash	ASTM D 3174, ASTM D 5142	Molybdenum	ASTM D 6357
Barium	INAA, ID-ICP-MS	Nickel	ASTM D 6357, ICP-MS
Beryllium	ASTM D 6357, ICP-AES ^{USGS}	Niobium	ICP-MS ^{USGS}
Bismuth	ICP-MS ^{USGS}	Nitrogen	ASTM D 3179, INA
Boron	PGAA	Potassium	ID-ICP-MS, INAA, PGAA
Bromine	INAA	Rubidium	ID-ICP-MS, INAA
Cadmium	ICP-MS	Samarium	INAA
Calcium	INAA	Scandium	INAA
Calorific Value	Commercial Coal Calorimeter	Selenium	ICP-MS, INAA
Carbon (Total)	ASTM D 3178, ICA	Silicon	WDXRF
Cerium	INAA	Silver	ICP-MS ^{USGS}
Cesium	INAA	Sodium	INAA, FES
Chlorine	ID-TIMS, PGAA, INAA	Strontium	ID-ICP-MS, INAA
Chromium	INAA	Sulfur	ID-TIMS
Cobalt	INAA, ICP-MS	Tellurium	ICP-MS ^{USGS}
Copper	ASTM D 6357, ICP-MS	Thallium	ICP-MS ^{USGS}
Europium	INAA	Thorium	ID-ICP-MS, INAA
Fluorine	ASTM D 5987-96, PI-GES	Tin	ICP-MS ^{USGS}
Gallium	ICP-MS ^{USGS}	Titanium	INAA
Germanium	ICP-MS ^{USGS}	Uranium	ID-ICP-MS
Hafnium	INAA	Vanadium	INAA
Hydrogen	PGAA, ASTM D 3178, IHA	Volatile Matter	ASTM D 3175, ASTM D 5142
Iron	INAA	Yttrium	ICP-AES ^{USGS}
Lead	ID-ICP-MS	Zinc	ICP-MS, INAA
Lithium	ICP-AES ^{USGS}	Zirconium	ICP-AES ^{USGS}

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Methods:

ASTM D 3178 Standard Test Methods for Carbon and Hydrogen in the Analysis Sample of Coal and Coke

ASTM D 3179 Standard Test Methods for Nitrogen in the Analysis Sample of Coal and Coke

ASTM D 5987-96 Standard Test Method for Total Fluorine in Coal and Coke by Pyrohydrolytic Extraction and

Ion Selective Electrode or Ion Chromatography Methods

ASTM D 6357 Standard Test Methods for Determination of Trace Elements in Coal, Coke, Combustion

Residues from Coal Utilization Processes by Inductively Coupled Plasma Atomic Emission, Inductively Coupled Plasma Mass, and Graphite Furnace Atomic Absorption Spectrometries

FES Flame Emission Spectrometry at NIST

FIA-CVAAS Flow Injection Cold Vapor Atomic Absorption Spectrometry at NIST

ICP-AES USGS Inductively Coupled Plasma Atomic Emission Spectrometry at United States Geological

Survey (USGS)

ICP-MS Inductively Coupled Plasma Mass Spectrometry at USGS ICP-MS Inductively Coupled Plasma Mass Spectrometry at NIST

ICA Instrumental Carbon Analyzer

ID-CV-ICP-MS Isotope Dilution Cold Vapor Inductively Coupled Plasma Mass Spectrometry at NIST

ID-ICP-MS Isotope Dilution Inductively Coupled Plasma Mass Spectrometry at NIST

ID-TIMS Isotope Dilution Thermal Ionization Mass Spectrometry at NIST

IHA Instrumental Hydrogen Analyzer INA Instrumental Nitrogen Analyzer

INAA Instrumental Neutron Activation Analysis at NIST
PGAA Prompt Gamma Activation Analysis at NIST
PI-GES Proton-Induced Gamma Emission Spectrometry
RNAA Radiochemical Neutron Activation Analysis at NIST
WDXRF Wavelength Dispersive X-ray Fluorescence at NIST

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Certificate Revision History: 31 January 2006 (This revision reflects a correction of the mass fraction unit of the ash content in Table 2 and editorial changes); 30 March 2004 (This revision reflects the addition of a fluorine reference value); 11 January 2001 (Original certificate date).

Users of this SRM should ensure that the certificate in their possession is current. This can be accomplished by contacting the SRM Program at: telephone (301) 975-6776; fax (301) 926-4751; e-mail srminfo@nist.gov; or via the Internet at http://www.nist.gov/srm.

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Addendum

Standard Reference Material® 1632c

Trace Elements in Coal (Bituminous)

CANSPECS 54 Coal Round Robin Results: SRM 1632c was included as an unknown in the February 1998 CANSPECS 54 Coal Round Robin. In addition, SRM 1632c coal was ashed in accordance to ASTM D 6357-00 [8] and included as an unknown ash in the February 1998 CANSPECS 54 Ash Round Robin. Summary statistics reported by CANSPECS are provided in the addendum to this certificate to demonstrate user experience with this material using conventional methods and to better characterize the matrix. The CANSPECS 54 Coal and Ash Round Robin results should **NOT** be used as substitutes for the NIST values.

Gaithersburg, MD 20899 Robert L. Watters, Jr., Chief Issue Date: 31 January 2006 Measurement Services Division

Addendum Revision History: 31 January 2006 (This revision reflects editorial changes); 30 March 2004 (Original addendum date).

See Addendum Revision History

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Summary of Analysis Reported by CANSPECS CANSPECS 54 Coal Sample NIST SRM 1632c

Parameter	Consensus Value	ASTM Method	ASTM Reproducibility	CANSPECS Reproducibility	ASTM Repeatability Standard Deviation	CANSPECS Repeatability	Number of Labs	Number of Methods
Moisture wt %	2.08	ASTM D 3173	0.11	0.13	0.07	0.06	79	22
Ash wt % db	7.16	ASTM D 3174	0.18	0.07	0.11	0.04	79	22
Volatiles wt % db	36.25	ASTM D 3175	0.35	0.87	0.18	0.19	65	17
BTU/lb db	13802	ASTM D 5865	44	64	18	18	73	16
Carbon wt % db	77.68	ASTM D 5373	0.89	0.78	0.23	0.22	33	14
Hydrogen wt % db	5.09	ASTM D 5373	0.11	0.20	0.06	0.07	32	13
Nitrogen wt % db	1.54	ASTM D 5373	0.06	0.07	0.04	0.03	34	13
Sulfur wt % db	1.49	ASTM D 4239c	0.05	0.05	0.03	0.02	77	18
Pyritic Sulfur wt % db	0.54	ASTM D 2492	0.10	0.12	0.05	0.02	16	4
Sulfate Sulfur wt % db	0.09	ASTM D 2492	0.01	0.05	0.01	0.01	16	3
Chlorine μg/g db	1127	ASTM D 4208	163	108	69	35	30	10
Fluorine µg/g db	70	ASTM D 3761	5	15	5	2	12	6
Mercury ng/g db	82	ASTM D 3684	11	13	7	8	15	8
Selenium μg/g db	1.32	ASTM D 4606	0.18	0.11	0.13	0.09	10	7
Free Swelling Index (FSI)	7.0	ASTM D 720	0.7	0.4	0.4	0.2	31	4

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Summary of Analysis Reported by CANSPECS CANSPECS 54 NIST SRM 1632c Laboratory Ash

		CHINDII	JCD 5+ 1115 1	DIGNI 10320	Edoordtory 1	1511		
Parameter	Consensus Value (in ash)	ASTM Method Referenced for Reproducibility and Repeatability	ASTM Reproducibility Standard Deviation	CANSPECS Reproducibility Standard Deviation	ASTM Repeatability Standard Deviation	CANSPECS Repeatability Standard Deviation	Number of Labs	Number of Methods
Al ₂ O ₃ wt %	23.91	ASTM D 4326	1.03	0.74	0.29	0.24	34	11
Antimony mg/kg	6.0	ASTM D 6357	1.0	0.4	0.6	0.3	10	4
Arsenic mg/kg	78	ASTM D 6357	7	15	4	3	13	6
BaO wt %	0.060	ASTM D 4326	0.026	0.006	0.013	0.004	19	6
Beryllium mg/kg	13.5	ASTM D 6357	1.5	2.4	0.4	0.4	9	6
Cadmium mg/kg	1.1	ASTM D 6357	0.2	0.3	0.07	0.06	9	6
CaO wt %	2.76	ASTM D 4326	0.15	0.10	0.07	0.06	34	12
Chromium mg/kg	189	ASTM D 6357	13	22	6	3	16	9
Cobalt mg/kg	41	ASTM D 6357	3	8	2	0.7	11	7
Copper mg/kg	82	ASTM D 6357	8	7	3	3	14	8
Fe ₂ O ₃ wt %	14.65	ASTM D 4326	0.53	0.50	0.10	0.17	34	11
K ₂ O wt %	1.75	ASTM D 4326	0.06	0.10	0.05	0.03	33	11
Lead mg/kg	46	ASTM D 6357	5	12	3	2	12	8
Manganese mg/kg	181	ASTM D 6357	13	12	7	3	15	8
MgO wt %	0.78	ASTM D 4326	0.07	0.03	0.04	0.02	32	11
Na ₂ O wt %	0.56	ASTM D 4326	0.15	0.06	0.07	0.02	33	12
Nickel mg/kg	126	ASTM D 6357	9	31	6	4	13	8
P ₂ O ₅ wt %	0.466	ASTM D 4326	0.08	0.064	0.024	0.016	29	8
SiO ₂ wt %	49.47	ASTM D 4326	1.56	1.31	0.45	0.28	33	11
SO ₃ wt %	2.73	ASTM D 4326	0.47	0.28	0.16	0.03	31	12
SrO wt %	0.099	ASTM D 4326	0.050	0.008	0.014	0.003	17	7
TiO ₂ wt %	1.09	ASTM D 4326	0.09	0.04	0.02	0.01	32	11
Vanadium mg/kg	324	ASTM D 6357	25	30	15	5	14	9
Zinc mg/kg	163	ASTM D 6357	11	31	6	4	14	9
Oxidizing Initial °C	1400	ASTM D 1857	20	9	11	4	20	1
Oxidizing Spherical °C	1435	ASTM D 1857	20	16	11	7	20	1
Oxidizing Hemi-spherical °C	1451	ASTM D 1857	20	11	11	6	20	1
Oxidizing Fluid °C	1471	ASTM D 1857	20	16	11	5	20	1
Reducing Initial °C	1193	ASTM D 1857	25	8	11	7	22	1
Reducing Spherical °C	1305	ASTM D 1857	20	43	11	6	22	1
Reducing Hemi-spherical °C	1338	ASTM D 1857	20	28	11	6	22	1
Reducing Fluid °C	1387	ASTM D 1857	30	27	11	7	22	1

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